



**Bihar Medical Services & Infrastructure Corporation
Limited**
2nd & 3rd Floor, Swasthya Bhawan, Behind IGIMS,
Sheikhpura, Adjacent to State Health Society, Bihar,
Patna- 800014
Phone/Fax: +919006405255, + 919471009193

CORRIGENDUM-I

Re-Tender for rate contract and supply of Quadrivalent Human Papilloma Virus (Serotypes 6, 11, 16 & 18) Vaccine (Recombinant) (qHPV) for different healthcare facilities of state of Bihar

Notice Inviting Tender Ref No.: -BMSIC/DRUGS/24-12/RETENDER dated: - 02-12-2024

(Only through E- Tender on website: -www.eproc2.bihar.gov.in)

Bihar Medical Services and Infrastructure Corporation Limited (BMSICL) invites E-Bids from the interested parties for “**Re-Tender for rate contract and supply of Quadrivalent Human Papilloma Virus (Serotypes 6, 11, 16 & 18) Vaccine (Recombinant) (qHPV) for different healthcare facilities of state of Bihar**”, vide Notice Inviting Tender No.- **BMSIC/DRUGS/24-12/RETENDER**. Detailed tender document containing eligibility criteria, selection mechanism, other terms and conditions are available on the website www.eproc2.bihar.gov.in.

After considering the suggestion/queries received from the prospective bidders & after due deliberation on all aspects, certain amendments have been made in the pack size as per Annexure- I and Clause 3(f) of Standard Bid Document as per Annexure- A in this Corrigendum-I.

**Sd/-
GM (Procurement)
BMSICL**

Annexure-I

S.N.	Name of Drug	Specification / Strength	Dosage Form	Pack Size	Amended Pack Size
1	Quadrivalent Human Papilloma Virus (Serotypes 6, 11, 16 & 18) Vaccine (Recombinant) (qHPV)	0.5 ml - 1 dose	Injection	0.5 ml Vial	0.5 ml Vial/PFS

Annexure A

Amendments in Bid Clause		
S.N.	Present Clause	Amended Clause
1	Clause 3(f)- • Minimum three years old valid Manufacturing License of the quoted product. • Manufacturing License along with approved product list must be valid till the last date of the submission of tender. • Market standing certificate & Manufacturing certificate issued by the Licensing Authority as a Manufacturer for each drug quoted for the last 3 years (Certificate should be enclosed with list of items) except for the drugs falling under the category of 'New Drug' as defined by CDSCO (Central Drugs Standard Control Organisation). If permission in Form 46 from DCGI has been obtained, then the 3 years Manufacturing & Market standing clause will be relaxed. The provisions of Rule 122 E of Drugs and Cosmetics Act rule 1945 shall be applicable. Explanation- In case of Importers Permission in Form 45 from DCGI is required as Per Drugs & Cosmetics Act 1940 & Rules 1945.	Clause 3(f) may be read as follows: - • Minimum one years old valid Manufacturing License of the quoted product. • Manufacturing License along with approved product list must be valid till the last date of the submission of tender. • Market standing certificate & Manufacturing certificate issued by the Licensing Authority as a Manufacturer for each drug quoted for the last 1 year (Certificate should be enclosed with list of items) except for the drugs falling under the category of 'New Drug' as defined by CDSCO (Central Drugs Standard Control Organisation). If permission in Form 46 from DCGI has been obtained, then the 3 years Manufacturing & Market standing clause will be relaxed. The provisions of Rule 122 E of Drugs and Cosmetics Act rule 1945 shall be applicable. Explanation- In case of Importers Permission in Form 45 from DCGI is required as Per Drugs & Cosmetics Act 1940 & Rules 1945.

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